PATENT COOPERATION TREATY

PCT

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Article 36 and Rule 70)

NO 7339WO FOR FURTHE			FOR FURTHER A	ACTION See Notification of Transmittal of International Preliminary Examination Report (Form PCT/IPEA/416)				
				International filing date	(day/mon	hiyear)	Priority date (day/mont/ 17.12.2002	uyear)
			ent Classification (IPC) or b 3L1/29	oth national classification	and IPC			
	STEC	S.A	. et al.					
1.			national preliminary exar and is transmitted to the				national Preliminary E	xamining
This REPORT consists of a total of 5 sheets, including this cover sheet.								
This report is also accompanied by ANNEXES, i.e. sheets of the description, claims and/or drawings which been amended and are the basis for this report and/or sheets containing rectifications made before this Aut (see Bulg 70.16 and Section 907 of the Administrative instructions under the PCTI.					ngs which have re this Authority			
	The	se an	nexes consist of a total c	of sheets.				
This report contains indications relating to the following Items:								
	I ☑ Basis of the opinion II □ Priority							
	- III	⊠	Non-establishment of o	opinion with regard to n	ovelty, in	ventive step ar	nd industrial applicabilit	ty
	IV Lack of unity of invention		on					
V A Reasoned statement under Rule 66.2(a)(ii) wi citations and explanations supporting such sta			ith regard atement	to novelty, inv	entive step or industria	al applicability;		
	VI Certain documents cited VII Certain defects in the international application		ed					
			1					
VIII Certain observations on the international application								
Date	of sub	missio	on of the demand		Date of c	completion of this	report	
09.0	07.20	04			03.05.2	2005		
			g address of the international	al	Authoriza	ed Officer	***************************************	Section Principles
	<u>a</u>))	D-8 Tel	ropean Patent Office 30298 Munich I. +49 89 2399 - 0 Tx: 52365	56 epmu d	Vernier	, F		
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 With regard to the elements of the international application (Replacement sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filled" and are not annexed to this report since they do not contain amendments (Rules 70.16 and 70.17).

	Des	scription, Pages			
	1-2		as originally filed		
	Claims, Numbers				
	1-1	1	as originally filed		
2.	. With regard to the language, all the elements marked above were available or furnished to this Authority in the language in which the international application was filed, unless otherwise indicated under this item.				
	These elements were available or furnished to this Authority in the following language: , which is:				
	the language of a translation furnished for the purposes of the international search (under Rule 23.1(b)).				
the language of publication of the international application (under Rule 48.3(b)).			ication of the international application (under Rule 48.3(b)).		
		the language of a tra Rule 55.2 and/or 55.3	nstation furnished for the purposes of international preliminary examination (under 3).		
3.	With regard to any nucleotide and/or amino acid sequence disclosed in the international application, the international preliminary examination was carried out on the basis of the sequence listing:				
		contained in the inter	mational application in written form.		
		filed together with the	e international application in computer readable form.		
		furnished subsequer	atly to this Authority in written form.		
		furnished subsequer	tty to this Authority in computer readable form.		
		The statement that the in the international a	ne subsequently furnished written sequence listing does not go beyond the disclosure oplication as filed has been furnished.		
		The statement that the listing has been furnit	ne information recorded in computer readable form is identical to the written sequence shed.		
4.	The	amendments have re	esulted in the cancellation of:		
		the description,	pages:		
		the claims,	Nos.:		
		the drawings,	sheets:		
5.			established as if (some of) the amendments had not been made, since they have to beyond the disclosure as filed (Rule 70.2(c)).		
		(Any replacement sh report.)	eet containing such amendments must be referred to under item 1 and annexed to this		
6.	Add	litional observations, i	f necessary:		

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Ш	III. Non-establishment of opinion with regard to novelty, inventive step and industrial applicability							
1.	The questions whether the claimed invention appears to be novel, to involve an inventive step (to be non-obvious), or to be industrially applicable have not been examined in respect of:							
	☐ the entire international application			on,				
	⊠ claims Nos. 11							
	because:							
	Ø	the said international application, or the said claims Nos. 11 relate to the following subject matter which does not require an international preliminary examination (specify):						
		see separate sheet						
		the description, claims or drawings (indicate particular elements below) or said claims Nos. are so unclear that no meaningful opinion could be formed (specify):						
		the claims, or said claims Nos. are so inadequately supported by the description that no meaningful opinior could be formed.						
		no international search report has been established for the said claims Nos.						
2.	ora	meaningful international preliminary examination cannot be carried out due to the failure of the nucleotide and ramino acid sequence listing to comply with the standard provided for in Annex C of the Administrative structions:						
		the written form has not been	furnist	ed or does i	not comply with the Standard.			
		the computer readable form h	as not	been furnish	ed or does not comply with the Standard.			
٧.	V. Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement							
1.	Stat	tement						
	Novelty (N)		Yes: No:	Claims Claims	1-11			
	Industrial applicability (IA) Yes: Claims			Claims Claims	1-11			
			Claims Claims	1-10				
2.	Cita	tions and explanations						
	see separate sheet							

EXAMINATION REPORT - SEPARATE SHEET

Item III

Claim 11 relates to subject-matter considered by this Authority to be covered by the provisions of Rule 67.1(iv) PCT. Consequently, no opinion will be formulated with respect to the industrial applicability of the subject-matter of this claim (Article 34(4)(a)(i)) PCT).

Item V

- The following documents are considered:
- D1: WO 96/31130 A (ABBOTT LAB) 10 October 1996 (1996-10-10)
- D2: US-A-4 303 692 (GAULL GERALD E) 1 December 1981 (1981-12-01)
- D3: DATABASE BIOSIS [Online] BIOSCIENCES INFORMATION SERVICE, PHILADELPHIA, PA, US; 1989, BRUNSER O ET AL: "EFFECT OF AN ACIDIFIED MILK ON DIARRHEA AND THE CARRIER STATE IN INFANTS OF LOW SOCIO-ECONOMIC STRATUM" XP002230476 Database accession no. PREV198988041353
- D4: ANONYMOUS: "17th International Congress of Nutrition" INTERNET ARTICLE, [Online] XP002230474 Retrieved from the Internet: URL:http://www.univie.ac.at/iuns2001/sw_3. htm> [retrieved on 2003-02-10]
- D5: ANONYMOUS: "Lactobacillus" INTERNET ARTICLE, [Online] XP002230475
 Retrieved from the Internet: URL:http://www.flora-balance.com/lactic_ac
 id_producing_bacteria.htm> [retrieved on 2003-02-10]
- The subject-matter of present independant claims does not meet the novelty requirements (Article 33(2) PCT) in the light of the disclosure in D1 (nutritional compositions acidified with lactic acid, see passages cited in the search report).
- Discussion of the question whether the claimed subject-matter involves an inventive step (Article 33(3) PCT) is only of relevance once novelty has been established.
- 4. The subject-matter of present claims 1-10 meets the requirements of Article 33(4) PCT, since it is applicable in the food industry. For the assessment of the present claim 11 on the question whether it is industrially applicable, no unified criteria exist in the PCT Contracting States. The patentability can also be dependent upon the formulation of the claims. The EPO, for example, does not recognize as industrially

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applicable the subject-matter of claims to the use of a compound in medical treatment, but may allow, however, claims to a known compound for first use in medical treatment and the use of such a compound for the manufacture of a medicament for a new medical treatment.